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(54) Title: CERAMIC FUSED FIBER ENHANCED DENTAL MATERIAL			
(57) Abstract <p>The preferred embodiment of the present invention provides novel and unique filler compositions and ceramic enhanced dental materials. The preferred embodiment of the composition and the ceramic dental restorative materials made therefrom comprise a rigid three-dimensional network of fibers fused together at their points of contact wherein said network is greater than 60 % by volume void space, has mean pore diameters greater than 10 microns, or both.</p>			

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TITLE OF THE INVENTION

Ceramic Fused Fiber Enhanced Dental Material

FIELD OF THE INVENTION

5 This invention relates to dental restorative materials. In another respect, the invention relates to ceramics and ceramic composite restorative materials. Yet in another aspect, the invention relates to novel restorative compositions, including but not limited to, silver-based mercury amalgams, castable ceramic/porcelain compositions, ceramic/metal hybrids, and plastic-based denture or denture-like materials for the direct or indirect
10 restoration or replacement of teeth or other oral anatomical structures. In a further aspect, the invention relates to novel restorative compositions for the direct filling of posterior teeth.

BACKGROUND OF THE INVENTION

15 Dental restorative materials include materials used to repair damaged teeth and/or replace missing teeth and/or other related oral structures. In some instances, dental restorative materials include materials used to reconstruct the maxillofacial complex. In general, dental restorative compositions include: (1) dental metal based amalgams, (2) enamel and dentin bonding agents, (3) dental hybrid composites, (4) dental cements and
20 bases, (5) casting alloys for crowns and bridges and other prosthetic structures, (6) ceramic/metal materials, (7) denture and prosthetic devices, (8) filled polymeric resins, (9) ceramic-based restorative materials, (10) impression materials, (11) sealants, and (12) temporary restorations and crowns, etc. The present invention relates to dental hybrid composite resins, denture and prosthetic materials, dental metallic based amalgams, tempo-
25 rary restorations and crowns, sealants, bases, cements, bonding agents, and ceramic porcelain restorative materials.

 A variety of compositions have been proposed and used for the direct filling of teeth. Of these compositions, some may be generally classified as dental composites and more specifically as resin composites. These resin composites are comprised of inorganic
30 particulates, i.e., filler, bound together with a polymeric matrix, i.e., a binder. The particulate filler reinforces the polymeric matrix and offsets its deficiencies. The binder, and/or polymeric matrix, may be comprised of an acrylic or epoxy resin or other types of carbon-based polymers. See, for example, U.S. Patent Nos. 3,066,112 and 3,179,623

which are hereby incorporated by reference. Fillers for such composite compositions, both posterior and/or anterior dental use, include finely divided solids like fumed silica, glass, zirconium, aluminum oxide, crystalline quartz, glass beads, or a mixture of glass beads and quartz or mixtures of the above materials. A material acceptable, however, for posterior use must be able to achieve a high filler loading capacity in the resin system. Moreover, filler strength, content, shape and size directly determines the physical and mechanical properties of the restoration material.

To date, there has been no composite material developed that completely meets the expected parameters needed for the intended use as a posterior dental restorative material to replace mercury-based dental amalgams. Dental materials presently available lack several physical or mechanical properties necessary for an ideal posterior dental restoration. As noted, it is imperative to achieve a high filler loading capacity in the resin system and presently all attempts to achieve such have failed. For example, highly loaded materials such as Microfine Composite™, using colloidal silica of a 40 nm size result in dramatically increased viscosity which jeopardizes handling characteristics. (See, Lambrechts, P.; Vanherle, G. (1983); *Structural Evidence of Microfilled Composites*. J. Biomed Mater Res 17:249-60; Willems, G.; Lambrechts, P.; Braen, M.; Celis, J.P.; Vanherle, G. (1993): *A Classification of Dental Composites according to their Morphology and Mechanical Characteristics*. Dent Mater 8:310-19). The colloidal silica forms an extended network structure that produces an increase in viscosity thereby limiting the amount of filler that can be incorporated to around 50% by volume. This 50% volume of filler loading has only been obtained by first filling to a higher degree, that is, greater than 50% during manufacturing, and then curing under high temperature and grinding to make colloidal oxide filled resin particles (organic fillers). However, a major problem still remains. The interface between these particles and the matrix, i.e., binder, is weak and causes brittle failure and wear. The filler composition of the present invention has the characteristics needed for posterior composite materials when combined with a resin matrix to address and solve these major hurdles.

The properties needed for an advantageous dental restorative material include, *inter alia*, the following: (1) low to high density, (2) high tensile/compression strength, (3) low thermal conductivity, (4) purity, (5) long life in cyclic applications, (6) high flexural strength, (7) rigidity, (8) inertness, (9) dimensional stability, (10) thermal shock resistance, (11) high diffusivity, (12) biocompatibility and (13) porosity. The present

invention provides heretofore unknown fused-fibrous dental restorative materials with the above properties.

SUMMARY OF THE INVENTION

5 An object of the present invention is to provide fused-fibrous/highly porous filler materials useful in dental fillings and materials that are particularly useful in dental restorative compositions, prosthodontic applications, and ceramic restorative materials, which have improved handling characteristics, improved strength, improved wear resistance, and decreased shrinkage from polymerization as compared to presently available
10 dental restorative compositions, dental prosthetic devices, and ceramic restorative materials. The composition of the present invention also contains less than 1% cristobalite contamination. It has been discovered that significantly improved particulates that are useful as fillers in dental restorative and prosthetic compositions are obtained from fused-fibrous compounds manufactured from: (1) from about 1% to about 50% by weight
15 alumina; (2) from about 50% to about 98% by weight silica; and (3) from about 1% to about 5% by weight boron. In addition, the composition may further comprise silicon carbide up to about 3% by weight. Also, the present invention is directed to a filler/reinforcer which is comprised of over 99% silica. The invention includes methods for the direct filling of teeth utilizing the filler/reinforcer of the present invention with
20 an initially liquid settable filling material comprising, in addition to filler, a binder wherein the material is allowed to harden after insertion in a tooth. In addition, the present invention provides reinforced dental restorative materials comprising a rigid three-dimensional network of inorganic or organic fibers fused together at their points of contact wherein said network has mean pore diameters of greater than 10 microns, or
25 is 60% or greater by volume void space, or both.

BRIEF DESCRIPTION OF THE FIGURES

FIGURE 1 is a scanning electron micrograph of the material of the present invention at 6 lb/ft³.

30 FIGURE 2 is a scanning electron micrograph of the filler/reinforcer material of the present invention at 16 lb/ft³.

FIGURE 3 is a graphical presentation of PRIMM density vs. porosity.

FIGURE 4 is a graphical presentation of the parameters of P.R.I.M.M.TM vs. Lucitone 199 Denture Acrylic.

FIGURE 5 is a graphical presentation of the parameters of P.R.I.M.M.TM vs. Restolux Sp-4 Composite Resine (Chopped Fibers).

5 FIGURE 6 is a graphical presentation of the parameters of P.R.I.M.M.TM vs. Herculite XRV (Kerr) Composite Resin.

FIGURE 7 is a graphical presentation of the parameters of P.R.I.M.M.TM vs. Tytin Amalgam Alloy.

10 FIGURE 8 is a graphical presentation of the Ideal Modulus form at density 16- Particle size greater than 180 μm - with silica (Colloidal) and Silanated Resin.

FIGURE 9 is a graphical presentation of the Compressive strength vs. the mean particle size, wherein significant improvement is shown with the addition of Colloidal Silica and presilanation of the resin.

15 FIGURE 10 is a graphical presentation of the (RA) Surface Roughness vs. Mean Particle Size, which shows a decrease in roughness with the addition of the silica (Colloidal).

DESCRIPTION OF THE PREFERRED EMBODIMENTS

20 Generally, the filler components of the present invention has been described above in the Summary Of The Invention. However, several fused-fibrous filler compositions falling within the description set forth above are particularly preferred. Generally, the compositions of the present invention are manufactured utilizing alumina fibers, silica fibers, and from about 1% to about 5% boron, preferably boron nitride.

25 One preferred embodiment is comprised of a binder and a filler wherein said filler is comprised of about 1% to about 50% by weight alumina, from about 50% by weight to about 98% by weight silica, and boron. Another preferred embodiment is manufactured utilizing: (1) from about 15% to about 30% by weight alumina fiber; (2) from about 65% to about 85% by weight silica fiber; (3) from about 1% to about 3% by weight silicon carbide; and (4) from about 1% to about 5% by weight boron nitride.

30 Another more preferred fused-fibrous composition for the filler is manufactured from: (1) about 21% by weight alumina fiber; (2) about 74% by weight silica fiber; (3) about 2% by weight silicon carbide; and (4) about 2.85% by weight boron nitride. Preferably, the material of the present invention is manufactured utilizing alumina and

silica fibers in a ratio of 22:78 by weight. For strength, a preferred ratio of alumina to silica is 30:70 by weight.

Moreover, a preferred embodiment of the composition of the filler, or reinforcement, of the present invention comprises: a rigid three-dimensional network of inorganic
5 or organic fibers fused together at their points of contact wherein said network has a mean pore diameters of greater than about 10 microns, or has greater than about 60% by volume void space, or both.

The ceramic/resin hybrid restorative materials of the present invention have the advantageous properties outlined above needed for dental restorative composites and
10 prosthetic materials. The filler material alone, has exceptional physical, chemical, and mechanical properties which are imparted to varying degrees depending upon the application to dental restoratives manufactured utilizing the material, these properties include, *inter alia*: (1) low to high density - 4 lb./ft.³ to 62 lb./ft.³; (2) low thermal conductivity - e.g., for 12 lb./ft.³ density at 500°F conductivity = 0.61 Btu - in./ft.²; (3) purity -
15 predominately comprised of 99.7% pure silica fibers and 95.2% pure alumina fibers; (4) long life in cyclic applications - e.g., 12 lb./ft.³ density, does not degrade with cyclic exposure to 2600°F and can even withstand limited exposure to 2900°F; (5) rigidity - maintains shape and supports mechanical loads while providing thermal insulation (i.e., has a high compressive strength and tensile strength (MN/m²) (6) high flexural strength
20 - modulus of rupture for 4 lb./ft.³ to 62 lb./ft.³ densities ranges from 100 - 6200 lb./in.²; (7) inert - does not burn, softens at temperatures above 2700°F and melts at about 3100°F; (8) dimensional stability - has a low coefficient of thermal expansion and 0.4% linear shrinkage; (9) thermal shock resistance - can be heated to 2600°F and immediately immersed in cold water without damage; (10) high diffusivity - 12 lb./ft.³
25 to 62 lb./ft.³ ranges from 97% to 56%; (11) porosity - highly porous (over 60% by volume void space, preferably between about 60% and about 98% by volume void space, most preferably over 80% by volume void space) and offers minimal resistance to the passage of gases or liquids (mean pore diameters greater than 10 microns, generally between 10 and 25 microns, most preferably from about 20 to about 22
30 microns); (12) able to coat or bond to other materials (i.e., materials, plastics, metals, inorganics) with relative ease to enhance characteristics. In addition, the 16 lb/ft³ density filler material of the present invention has (1) a flexural modulus of strength - 2.5×10^{10} Pa, (2) a Rockwell Hardness - 50, (3) Surface Roughness of 0.6 Ra. and (4)

linear shrinkage of 0.4% after repeated cycles at temperatures above 2700°F. As a result, e.g., the composite resins made with the filler of the present invention have heretofore unknown linear shrinkages of less than about 0.09%.

Generally, the process for the manufacture of fused-fibrous silica/alumina and/or other ceramic fiber of low density, like 16 lb. per ft.³, is comprised of the following steps:

- (1) preparation of a slurry mixture comprised of pre-measured amounts of purified fibers/materials and deionized water;
- (2) removal of shot from slurry mixture;
- 10 (3) removal of water after thorough mixing to form a soft billet;
- (4) addition of a ceramic binder after the formation of the billet;
- (5) placement of the billet in a drying microwave oven for moisture removal; and
- 15 (6) sintering the dry billet in a large furnace at about 1600°F or above.

The high purity silica fibers above are first washed and dispersed in hydrochloric acid and/or deionized water or other solvents. The ratio of washing solution to fiber is between 30 to 150 parts liquid (pH 3 to 4) to 1 part fiber. Washing for 2 to 4 hours generally removes the surface chemical contamination and non-fibrous material (shot) which contributes to silica fiber devitrification. After washing, the fibers are rinsed 3 times at approximately the same liquid to fiber ratio for 10 to 15 minutes with deionized water. The pH is then about 6. Excess water is drained off leaving a ratio of 5 to 10 parts water to 1 part fiber. During this wash and all following procedures, great care must be taken to avoid contaminating the silica fibers. The use of polyethylene or stainless steel utensils and deionized water aids in avoiding such contamination. The washing procedure has little effect on the bulk chemical composition of the fiber. Its major function is the conditioning and dispersing of the silica fibers.

The alumina fibers are prepared by dispersing them in deionized water. They can be dispersed by mixing 10 to 40 parts water with 1 part fiber in a V-blender for 2 1/2 to 5 minutes. The time required is a function of the fiber length and diameter. In general, the larger the fiber, the more time required.

Generally, in order to manufacture ultra-low density fused-fibrous ceramic filler material, for example, densities below 12 lb./ft.³, the process includes the additional steps of:

- (1) the addition of expendable carbon fibers in the casting process and/or other temporary support material; and
- (2) firing the billet at about 1300°F to remove the carbon fibers or other support material prior to the final firing at approximately 1600°F or above.

When the dispersed silica fibers and dispersed alumina fibers are combined, the pH is probably acidic and should be adjusted to neutral with ammonium hydroxide. The slurry should contain about 12 to about 25 parts water to about 1 part fiber. The slurry is mixed to a uniform consistency in a V-blender in 5 to 20 minutes. The boron nitride can be added at this point (2.85% by weight of the fibers) and mixed to a uniform consistency in a V-blender for an additional 5 to 15 minutes creating a Master Slurry. The preferred mixing procedure uses 15 parts water to 1 part fiber and the slurry is produced in about 20 minutes of mixing. At lower density formulations, expendable carbon fibers are used to give "green" strength to the billet prior to the final sintering. The percent of carbon fiber used varies greatly depending on the diameter, length and source of the fiber and the ultimate density of the material being produced. The percent of carbon fiber per dry weight of material should range between 1 and 10%. The source of the carbon fiber can take many forms including nylon, cellulose, and purified graphite based carbon in fibrous form. Carbon fibers added in the casting process are eliminated by firing the billets at 1350°F prior to the final firing at 2450°F.

The Master Slurry is poured into a mold for pressing into the desired shape. The water is withdrawn rapidly and the resulting felt is compressed at 10 to 20 psi. Rapid removal of the water is required to prevent the fibers from separating. If graded properties are desired in the resultant material, the slurry can be allowed to settle and the fibers to partially separate before the removal of the water.

The final density of the finished restorative material is determined in part by the amount of compression placed on the felt, varying the wet molded dimension in relation to the fiber content. The formulation of the present invention has been prepared in densities ranging from about 0.05 to 0.48 g/cc. It can, however, be prepared in lower and higher densities.

After molding, the restorative material is dried and fired by the following preferred procedure. The material is first dried in an oven for 18 hours; the temperature, initially 38°C, is raised at a rate of 11°C per hour to 104°C, held there for 4 hours, raised again at a rate of 11°C per hour to 150°C, and held there for 4 hours.

5 The material is taken directly from the drying oven, placed in the firing furnace, and fired. A temperature rise rate of 220°C per hour or less is required in order to avoid cracking and warping in the case of a 15 cm x 15 cm x 7.5 cm block of material. For larger blocks, slower heating rates may be required. The maximum firing temperature may vary from 1200°C to 1600°C depending upon the fiber ratio used, amount of

10 boron nitride, and the final density of the material that is desired.

The temperature rise rate is chosen to permit relatively uniform temperatures to be achieved throughout the material during the process. A faster temperature rise rate causes non-uniform temperatures to be achieved throughout the material during the process. A faster temperature rise rate causes nonuniform strength and density and may

15 cause cracking. Longer or higher temperature firing results in higher shrinkage and related greater resistance to subsequent shrinkage, as well as a shorter lifetime to devitrification under cyclic exposures to high temperatures. The maximum firing temperature is dependent upon the fiber ratio used and the density of the composite desired. The firing time and maximum temperature are selected to allow sufficient

20 shrinkage to achieve stabilization and fiber fusion while not allowing any devitrification.

After firing, the material may be machined to obtain any desired final dimensions. Only about 0.5 cm of the material must be machined off.

The procedure used to prepare this restorative material, i.e., the polymeric rigid inorganic matrix material of the present invention, may be varied through a rather broad

25 range with satisfactory results. In one variation, the silica fibers may be borated and prefired prior to use. This process is used to improve the morphological stability and physical properties of the resultant material.

The following examples are provided to illustrate the invention by describing various embodiments, including its best mode as presently conceived. All proportions

30 used are expressed on a percent by weight basis unless otherwise noted.

EXAMPLE 1

An embodiment of the fused-fibrous matrix ceramic material of the present invention having a density of 0.32 g/cc, and opacified with silicon carbide was produced.

with 825 grams of silica fibers, 175 grams alumina fiber (average diameter - 11 microns, length - 0.32 cm), 35 grams 1200 grit silicon carbide, 2.85 grams of boron nitride, 10 milliliters hydrochloric acid, 5 milliliters ammonium hydroxide and deionized water. The silica fibers were washed as in Example 2.

5 The alumina fibers were placed in a 7,570 ml capacity stainless steel double shell blender with 5,000 grams deionized water and mixed using an intensifier bar for 2 1/2 minutes to disperse the fiber.

10 The washed silica fibers, dispersed alumina fibers, boron nitride, and silicon carbide were placed in a 28.31 liter stainless steel double shell V-blender. Deionized water was added to bring the total weight to 15,000 grams. The ammonium hydroxide (5 ml) was added to adjust the slurry to basic before mixing. The slurry was mixed, degassed, transferred to a mold and pressed into a billet as in Example 2.

EXAMPLE 2

15 The materials used were the following: 150 grams aluminasilicate fibers (AS-32, manufactured by 3-M Company containing 80% Al_2O_3 and 20% SiO_2), 1000 grams of silica fibers (Microquartz 108), 35 grams of 1200 grit silicon carbide, 30 grams of boron nitride, 10 ml of hydrochloric acid, 5 ml of ammonium hydroxide, and deionized water.

20 The silica fibers were placed in a polyethylene container in 32 liters of deionized water. Hydrochloric acid (10 ml) was added to bring the pH to 3. Pure nitrogen was bubbled through the mixture to agitate the fiber and assist washing. Agitation was continued for two hours. The acidic water was then drained off, fresh deionized water added and the mixture again agitated with pure nitrogen for 15 minutes. The rinsing process was repeated 2 more times which brought the pH to about 6.0.

25 The aluminasilicate fibers were placed in a 7,570 ml capacity stainless steel double shell blender with 5,000 grams of deionized water and mixed using the intensifier bar for 2 1/2 minutes to disperse the fiber.

30 The washed silica fibers, dispersed aluminasilicate fibers, boron nitride, and silicon carbide were placed in a 28.31 liter stainless steel double shell V-blender. Deionized water was added to bring the total weight to 18,000 grams. Ammonium hydroxide (5 ml) was added to adjust the slurry to basic before mixing. The slurry was then mixed with the intensifier bar running for 20 minutes, removed from the V-blender and degassed, transferred into a mold, and pressed into a 21.6 cm x 21.6 cm x 10 cm

billet. The top and bottom of the mold were perforated and covered with a 16 mesh aluminum screen to allow the excess water to flow out.

The billet was dried in an oven for 18 hours beginning at 38°C, increased at 11°C per hour to 104°C, held for four hours at 104°C, increased at 11°C per hour to 150°C and held four hours at 150°C. After drying, the billet was transferred to the firing furnace. The furnace temperature was increased at a rate of 220°C per hour to the firing temperature, 1315°C, where it was held for 1 1/2 hours. After firing, the temperature was decreased at a rate of 220°C per hour to 980°C where the furnace was turned off, then allowed to cool to room temperature.

The usefulness of boron oxide in the two-fiber composites of this invention is demonstrated by the following preparations.

EXAMPLE 3

In one run, an experimental mixture was made containing 25% aluminasilicate fibers ("FIBERFRAX H," manufactured by the Carborundum Company, containing 62% Al_2O_3 and 38% SiO_2) and 75% pure silica fibers ("MICROQUARTZ 108"). The mixture was ground with mortar and pestle and then fired at 1400°C for 5 hours. The resulting product underwent 48% devitrification. When the aluminasilicate fibers were prefired with boron oxide (85% and 15% respectively) at 1100°C for 90 minutes and then mixed with the silica fibers and fired as above, the product exhibited no devitrification.

EXAMPLE 4

An acceptable 17 cm x 17 cm x 7.5 cm billet of material having a density of 0.11 g/cc was produced using 600 grams of silica fibers, 90 grams of aluminaborosilicate fibers (average diameter-11 microns, 0.64 cm long), 10 ml of hydrochloric acid, 5 ml of ammonium hydroxide, and deionized water.

The silica fibers were washed in accordance with the procedure of Example 2. The aluminaborosilicate fibers were dispersed in a 7,570 ml V-blender with 3000 grams of deionized water for 5 minutes. The washed silica fibers, dispersed aluminaborosilicate fibers, and ammonium hydroxide were mixed, with enough deionized water to bring the total weight to 9,000 grams, in a 28.31 liter V-blender for 10 minutes with the intensifier bar running. The slurry was removed from the V-blender, degassed, molded and the resulting billet fired as in Example 2. The billet was then transferred to the firing furnace. The furnace temperature was increased at a rate of 220°C per hour to the firing temperature, 1260°C, where it was held for 5 hours. After firing, the temperature

was decreased at a rate of 220°C per hour to 980°C, at which point the furnace was turned off and allowed to cool at room temperature. The billet was machined to 17 cm x 17 cm x 7.5 cm in accordance with usual machining practices.

EXAMPLE 5

5 An acceptable 17 cm x 17 cm x 7.5 cm billet of material with yet greater stability toward devitrification than the material produced in example 1, having a density of 0.32 g/cc, and opacified with silicon carbide was produced using 825 grams of silica fibers, 175 grams aluminaborosilicate fibers (average diameter-11 microns, 0.64 cm long), 35 grams of 1200 grit silicon carbide, 10 ml of hydrochloric acid, 5 ml of ammonium
10 hydroxide, 56.8 grams of boron oxide, and deionized water.

 The silica fibers were washed in accordance with the procedure of Example 2. The boron oxide was dissolved in 4,000 grams of deionized water (concentration-1.42% boron oxide). The aluminaborosilicate fibers were placed in a stainless steel basket and dipped into the boron oxide solution (the aluminaborosilicate fibers absorbed 7 times
15 their own weight of the boron oxide solution). The fibers with absorbed boron oxide were then dried at 104°C for 4 hours and calcined at 1100°C for 1 hour. The "borated" fibers were then placed in a 7,570 ml capacity stainless steel V-blender with 5,000 grams of deionized water and mixed using the intensifier bar for 2 1/2 minutes to disperse the fiber. The washed silica fibers, dispersed "borated" aluminaborosilicate
20 fibers, silicon carbide, and ammonium hydroxide were mixed with enough deionized water to bring the total weight to 15,000 grams, in a one cubic foot V-blender for 20 minutes with the intensifier bar running. The slurry was removed from the V-blender, degassed, molded, dried, fired, and machined, as in Example 1.

EXAMPLE 6

25 An acceptable 17 cm x 17 cm x 7.5 cm billet of material with graded properties, having a density of 0.32 g/cc, and opacified with silicon carbide, was produced using 825 grams of silica fibers, 175 grams of aluminaborosilicate fibers (average diameter-11 microns, 0.64 cm long), 35 grams of 1200 grit silicon carbide, 10 ml of hydrochloric acid, 5 ml of ammonium hydroxide, and deionized water.

30 The silica fibers were washed in accordance with the procedure of Example 2. The aluminaborosilicate fibers were dispersed in a 7,570 ml V-blender with 5000 grams of deionized water for 5 minutes. The washed silica fibers, dispersed aluminaborosilicate fibers, silicon carbide and ammonium hydroxide were mixed with enough deionized water

to bring the total weight to 25,000 grams, in a 28.31 liter V-blender for 15 minutes with the intensifier bar running. The slurry was removed from the V-blender, degassed, molded, dried, fired and machined in accordance with the procedure of Example 1.

5 The resulting billet of material is relatively richer in silica at the top and aluminaborosilicate at the bottom.

EXAMPLE 7

A 17.5 cm x 17.5 cm x 9 cm material with a temperature capability greater than that of the material of Example 1, having a density of 0.24 g/cc, and opacified with silicon carbide, was produced using 750 grams of aluminaborosilicate fibers (diameter-1
10 to 3 microns), 250 grams of silica fibers, 35 grams of silicon carbide, 5 ml of ammonium hydroxide, and deionized water. The silica fibers were dispersed in a 7,570 ml V-blender with 5,000 grams of deionized water for 5 minutes.

The dispersed silica fibers, aluminaborosilicate fibers, silicon carbide, and ammonium hydroxide were mixed with enough deionized water to bring the total weight to
15 18,000 grams, in a 28.31 liter V-blender for 7 minutes with the intensifier bar running. The slurry was removed from the V-blender, degassed, molded, and dried as in Example 2. In the furnace, the temperature was increased at a rate of 220°C per hour to the firing temperature of 1370°C where it was held for 1 1/2 hours. After firing, the temperature was decreased at a rate of 220°C per hour to 980°C, at which point the
20 furnace was turned off and allowed to cool to room temperature. The billet was machined to 17.5 cm x 17.5 cm x 9 cm in accordance with the usual machining practices.

The preferred alumina fibers are 95.2% pure and are available from ICI Americas, Inc. and marketed as SAFFRIL™. The preferred diameter for the alumina fibers ranges
25 from 1 to about 15 microns. The preferred silica fibers are 99.7% pure and are available from Schuller (Johns Manville Corp.), Denver, Colorado and marketed as MICROQUARTZ 108™ fibers or as Q-FIBER™. These fibers have an average diameter of 1.7 microns. However, silica fibers having diameters ranging from 1 to 6 microns are useful in the present invention.

30 Also, mixtures of the above-described fibers can be used with other fibers known in the art, e.g., zirconium fibers. In addition, metal fibers and carbon fibers can be utilized by themselves or in combination with other fibers. As stated, the product of

the method of the present invention to make the filler/reinforcer of the present invention may comprise as much or greater than 99% silica.

While boron nitride is considered to be the preferred boron source, it is believed that SiB_x , B_4C , B_2O_3 , and B and other boron sources can also be used. It is preferred that boron be present in an amount from about 0.4% to about 3% by weight. Boron nitride is believed to be preferred because it is believed, due to its stability, that it permits a more uniform fusion to fiber junction and yields superior bonding and uniform porosity.

In addition, aluminaborosilicate fibers may be used and are available from 3M Company marketed under the tradename AB-312™ which contains 62% (plus/minus 2.0%) Al_2O_3 , 14% (plus/minus 2.0%) B_2O_3 and 24% (plus/minus 2.0%) SiO_2 . These fibers are available and useful in the present invention in diameters ranging from 3 to 12 microns.

The preferred composition comprised of: 21% by weight alumina fiber; 74% by weight silica fiber; 2% by weight (600 grit) silicon carbide; and 2.85% by weight boron nitride is also available commercially in densities to 3 to 64 lbs./ft.³ (plus/minus 3/4 lb.) from Lockheed Missiles and Space Company, Inc., Sunnyvale California ("Lockheed") under the trade name "HTP" (High Temperature Performance). For example, Lockheed commercially sells "HTP-12-22" (12 lb./ft.³ density silica/alumina fiber ratio of 78/22), "HTP-12-35" (12 lb./ft.³ density in a silica/alumina fiber ratio of 65/35) and "HTP-12-45" (12 lb./ft.³ density in silica/alumina fiber ratio of 55/45).

Materials Evolution & Development USA, Inc. manufactures the above types of Ultra-low density fused-fibrous ceramics under the trade name P.R.I.M.M.™ (Polymeric Rigid Inorganic Matrix Material). Materials Evolution & Development USA, Inc. manufactures P.R.I.M.M.™ in densities varying from 4 lb./ft.³ to 64 lb./ft.³. P.R.I.M.M.™ material is ground by mortar and pestle, or other grinding mechanism, then sieved into different particulate sizes. An optimum sieved particulate size for P.R.I.M.M.™ material (16 lb./ft.³ density) is approximately 180 microns in diameter. However, optimum particle sizes are believed to be dependent upon the surface tension and viscosity of the binder used in a particular application and P.R.I.M.M.™ material density and can range from about 150 microns to about 350 microns. The preferred particle size, i.e., the filler or reinforcement particles, of the PRIMM™ product of the present invention ranges from about 180 microns to about 250 microns. In addition, in

some applications it is preferable to utilize particles of a smaller size in combination with the above particle size range. For example, in composite resin systems, it is preferable to utilize 3 parts particulates in the 180-250 micron range to 1 part particulates in the 90-125 micron range.

5 In addition to the filler/reinforcer of the present invention, other fillers known in the art may be utilized in combination in binder systems, for example, microfil fillers selected from the group consisting of inorganic materials, metallic materials, organic materials, and mixtures thereof. Examples include barium glass, zirconium, crystalline carbon and mixtures thereof. Indeed, utilizing the present inventive filler/reinforcer with
10 microfills, a composite resin may comprise over 80% filler loading, preferably over 90% filler loading by weight.

The present invention includes the use of radiopacifiers including, but not limited to, TPB, bismuth, aluminum compounds, metal oxides, and organo-metallic compounds.

The present invention may be utilized with a variety of binder or resin systems
15 including, but not limited to, self-activating, light curable, heat curable, self-curing and microwave curing systems.

The density of the PRIMM™ product, i.e., the filler or reinforcer, of the present invention ranges from about 6 lb/ft³ to about 50 lb/ft³, preferably from about 6 lb/ft³ to about 25 lb/ft³, and most preferably from about 6 lb/ft³ to about 16 lb/ft³.

20 The porosity of the PRIMM™ product, i.e., the filler or reinforcement, of the present invention ranges from about 60% by volume void space and up, preferably the product porosity ranges from about 70% by volume void space and up. and most preferably, the porosity of the reinforcement of the present invention ranges from about 80% by volume void space. For example, 16 lb/ft³ PRIMM™ product of the present
25 invention manufactured utilizing the examples of the present invention comprises approximately 88% by volume void space. The relationship between density and porosity is shown in Figure 3.

The reinforcement or filler of the present invention (PRIMM™ product) preferably has average or mean pore diameters of greater than 10 microns, more preferably over
30 20 microns. For example, see FIG. 1, an SEM of an example of the PRIMM™ product of 16 lb/ft³ density, silanated as described herein, and with colloidal silica. As can be seen, the material comprises mean pore diameters greater than 10 microns.

The compressive strength (MN/m^2) and tensile strength (MN/m^2) of an embodiment of the restorative material of the present invention (78% alumina, 22% silica, % 3 silicon carbide, 2.85% boron nitride at 16 lb/ft^3 density) is compared in Figures 4-7 to commercially available dental restorative materials: "LACITONE 199 DENTURE ACRYLIC" available from Dentsply International, Inc.; "RESTOLUX SP-4 COMPOSITE RESIN" (Chopped Fibers) available from Lee Pharmaceuticals; "HERCULITE XRV" (Kerr) composite resin available from Kerr Manufacturing Co.; "TYTIN AMALGAM ALLOY" available from Kerr Manufacturing Co.

The tensile and compressive strength values of the commercially available materials used for comparison were taken from product information sheets provided by the manufacturers. The method used to determine the tensile strength of the material of the present invention was the Brazilian Test as described in the literature of Craig. (See, Craig, R.G. (1989); *Restorative Dental Materials*, 8 Ed., St. Louis: Mosby, 65-12; 255-92 which is hereby incorporated by reference.)

Figures 4-7 illustrate the compressive and tensile strength advantage of the material of the present invention over several commercially available materials. As shown and tested, the material of the present invention has a compressive strength equal to or greater than about 550 MN/m^2 and a tensile strength equal to or greater than about 80 MN/m^2 .

In relation to mixing the filler component of the present invention with a binder to form the composite of the present invention, approximately 1.0 gram of P.R.I.M.M.TM fused-fibrous ceramic product, having a density of 16 lb/ft^3 to 28 lb/ft^3 , was added to approximately 1 gram of GTE resin available from Dentsply and mixed until a complete wetting occurred. It should be noted, that this resin ratio was found to be ideal for easy incorporation and loading beyond this point was detrimental to the ultimate handling of the material. In fact, at a point past this ratio, the composite became very thick and dry and the overloading resulted in unusable material. It is also believed that an increased particle cluster size improves the capillary action of the fibers aligned for an easier incorporation of the resin into the fibers.

It has also been found to be preferable, that once the fibers have been thoroughly mixed into the resin, a known amount of colloidal silica (0.3 grams) ("CABOT-CABOSIL LM 130" fumed silica) was incorporated until the material had a consistency similar to that of currently marketed composite resins. However, unlike currently

available composite resins, this mixture proved to be packable and handled much the same way as an amalgam alloy.

It is also believed that the addition of a silanation agent improves the fiber to resin bond. For example, an organofunctional silanation agent (Union Carbide-Silane A-174) was added to the composite. Specifically, 0.1 gram of the silane was added to 1.0 gram of resin prior to mixing it with the fibers. The result of this action produced an intact fiber resin interface bond following the fracturing of the sample. The preferred procedure of silanation of fibers is as follows: (1) mix 4 ml silane (Union Carbide A-179) with 4 ml n-propylamine with 192 ml of cyclohexane for 15 minutes; (2) treat filler fibers for 2 hours in the above prepared solution and stir at room temperature; (3) rinse once with cyclohexane; (4) dry in air at room temperature for 1 hour; and (5) dry at 60° C for 1 hour. Preferably, the fibers are silanated greater than 90% of the surface area of the fibers.

The silanation process to produce a presilanated binder, e.g., polymer, requires the addition of 0.1% by weight of A-174 (Union Carbide Silane; gamma methacryloxy propyltrimethoxy silane) to the predetermined amount of binder needed to formulate the optional fiber-binder ratio prior to mixing.

In addition, it has been found to be preferable when adding the filler/reinforcer of the present invention to resin systems, to do so under vacuum pressure. Also, it is deemed preferable to add the filler/reinforcer of the present invention to resin systems at an elevated temperature, i.e., greater than 25° C to decrease the viscosity of the resin system.

Preferred organic binders for use with the filler or reinforcement of the present invention include, but are not limited to, acrylate resins, for posterior composite resins, dentures, sealants, laboratory resins, temporary crowns, and denture teeth. For plastic restoration, epoxy resins are preferred.

Preferred inorganic binder for use with the filler/reinforcement of the present invention include, but are not limited to, zinc oxide for bases, zinc phosphate and glass ionomers for cements, 4-meta for bonding agents, calcium hydroxide for liners, and lucite for porcelains.

Preferred metallic binders include, but are not limited to, for use with the filler/reinforcers of the present invention: silver/copper/tin/mercury amalgam, gallium, gold, noble metal and titanium.

It has also been found that the mean particle size (microns) effects the flexural modulus ($\text{Pa} \times 10^{10}$), compressive strength (MN/m^2), and surface roughness (Ra). Shown in Figures 8-10, is the filler component of the present invention at 16 lb/ft³ density (1:1 filler/resin mixture) for different particle sized (microns). Note, as used in Figures 8-10, the particle sizes designated with a double asterisk "**" refer to material which includes colloidal silica and the particle sizes designated with a single asterisk "*" refer to material which includes colloidal silica wherein the resin was silanated.

A comparison of the properties of the filler component of the present invention as compared to prior art filler components is shown in Table 1.

TABLE 1

Product	P.R.I.M.M. TM FCC + SiO ₂ *	Restolux SP- 4 TM	Z100 (3M)	XRV Herco.	TPH Prisma	Heliomolar	Charisma
Filler	Alumi- na/Silica (Fused Fi- bers)	Chopped Glass Fibers	Zirconia/ Silica	Barium Glass	Barium Glass	SiO ₂	Barium Glass/ Microglass
Filler loading Wt (%)	60	68.5	66	56	53	38	73
Average Particle Size (microns)	180/0.4	300	0.6	0.6	1	Microfil** (hybrid)	0.7
Diametral Tensile (MPa)	82	**	83	81	72	47	77
Compressive Strength (MPa)	560	300	448	324	375	436	399
Hardness (Kg/mm ²)	50	125	78.3	53.3	43.5	35.9	**
Y-Modulus (Pa)	25×10^9	23×10^9	13×10^9	7.6×10^9	5.7×10^9	4.3×10^9	++
Surface Roughness (Ra)	0.30	1.19	0.27	0.12	0.29	.013	**
Coeff of Thermal Expansion $\times 10^6$ unit/unit/C	10	17	17	27	30	52	38
Radiopacity	TPB***[Yes]	Yes	Yes	Yes	Yes	Yes	Yes
Linear Shrinkage	6.08	0.35	0.29	**	.03	**	**

* PRIMMTM (1.0g) containing FCC's and colloidal silica of 0.4mm diameter with 1.0g Dentsply GTE resin (Bis GMA, Bis EMA, Teg DMA).

** Data unavailable.

*** "TPB" - Radiopaque additive of proprietary nature, available from Sluka of Germany having the formula C₈H₁₅Bi, equivalent to 16mm of aluminum.

TABLE 2
PRIMM™ AUGMENTED AMALGAM

		KN	MPa
	Control	1250	99.2
5	(Dispersalloy®)	1275	101.2
		1100	87.3
		1000	79.4
		1000	79.4
		1230	97.6
10	2% PRIMM	2500	198.4
		2225	176.6
		1850	146.8
		1850	146.8
		1600	127.0
15		1750	138.9
	4% PRIMM	2050	165.9
		1750	138.9
		1850	146.8
		1800	142.9
20		1650	131.0
		1600	127.0

The above results report on incorporation of PRIMM™ (16 lb/ft³) into Dispersalloy® available from Dentsply Caulk. In the above Table, three groups are reported on: (1) 100% Dispersalloy® available from Dentsply Caulk. In the above Table, three groups are reported on: (1) 100% Dispersalloy® (no PRIMM™ product), (2) 98% by weight Dispersalloy® and 2% by weight PRIMM™ product, and (3) 96% by weight Dispersalloy® and 4% by weight PRIMM. The fibers of the PRIMM™ product were presilanated as discussed above. They were then treated with "All-BOND II" available from Bisco, Inc. "All-BOND II" is known as a 4-META adhesive comprising the adhesive molecule 4-methacryloxyethyl trimellitate anhydride. The test specimens were made using 9mm long polypropylene tubes with an inner diameter of 4mm. The

control specimens were prepared on glass plates and packed into the tubes using standard dentistry procedures. For the augmented specimens, the PRIMM™ product particles were mixed into the amalgam to a uniform consistency and the blend packed into the tubes. After curing, the plastic tubing was removed and the specimens milled to a length of 8mm. The specimens were tested in an Instron Universal testing machine (available from Instron Corp.). The cylinders were placed in a steel holder which aligned the top surface parallel with the flat surface of the crosshead to insure even distribution of the applied force. The cylinders were subjected to continuous loading at a crosshead speed of 5mm/min. until fracture. The compressive forces required for fracture were then recorded in Newtons and calculated in Megapascals (Mpm) using the formula:

Fracture force: Newtons

Cylinder surface area: radius = 2mm; area = $\pi(2)^2 = 12.6\text{mm}^2$

Compressive Strength: Fracture force/cylinder surface area =
Newtons/mm² = Megapascals.

While the invention has been illustrated and described in detail in the foregoing description, the same is to be considered as illustrative and not restrictive of character, it being understood that only the preferred embodiment has been shown and described and that all changes and modifications that come within the spirit of the invention are desired to be protected.

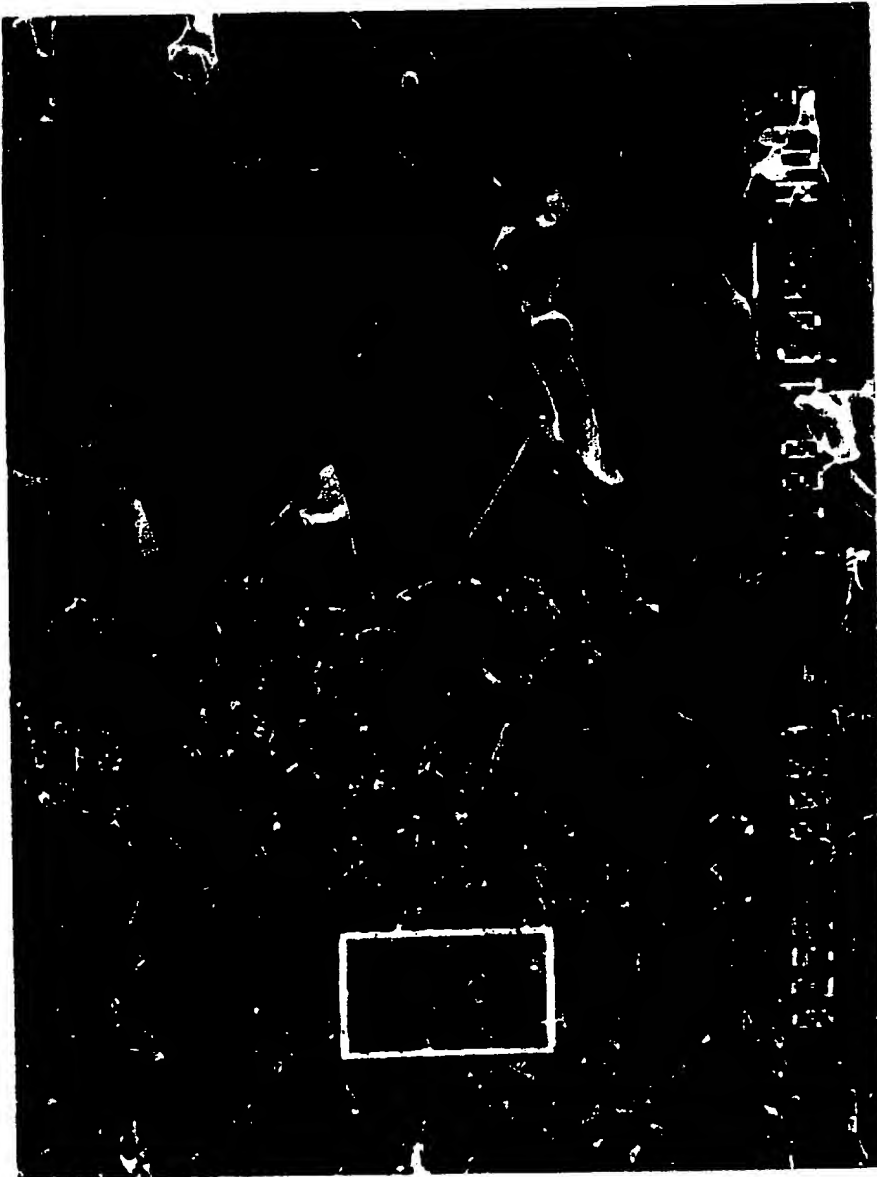
What is claimed is:

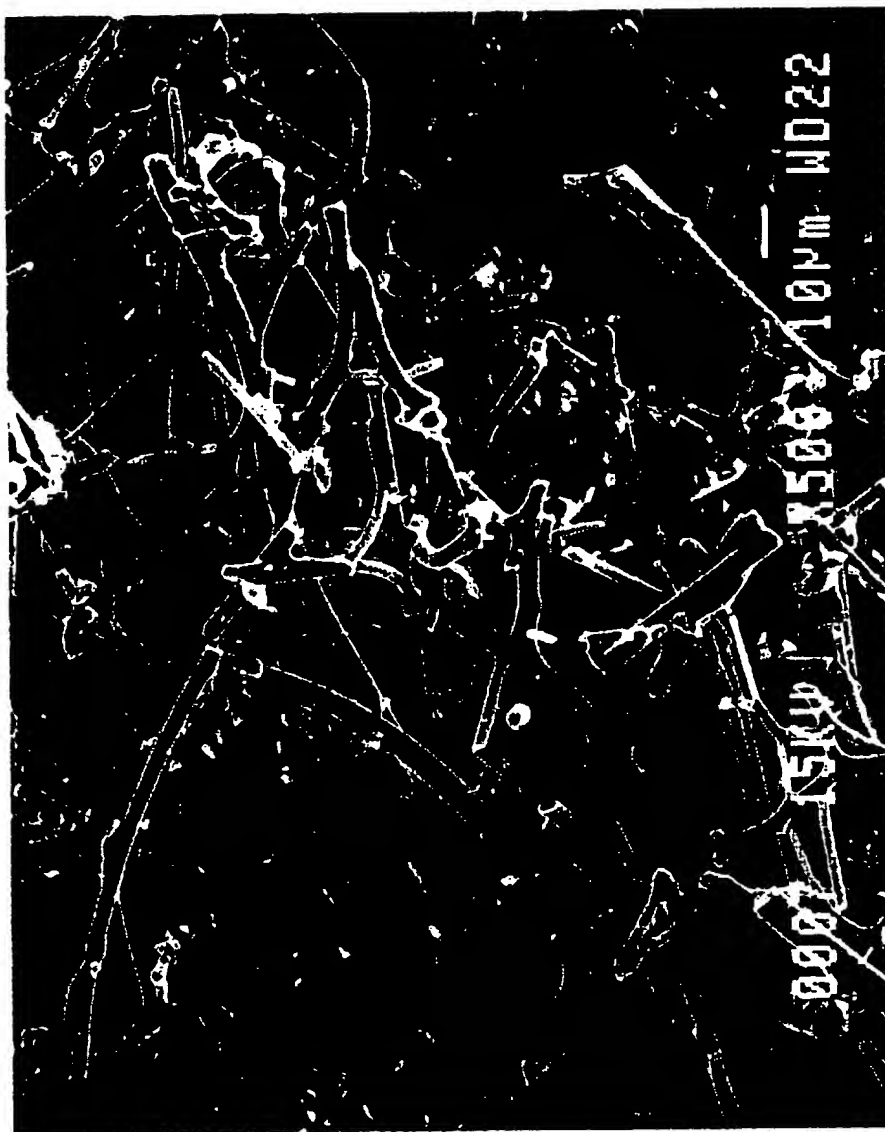
- 1 1. In a method for the direct filling of a cavity in a tooth, wherein said method
2 includes filling said cavity with an initially liquid, settable filling material comprising a
3 filler and a binder admixed therewith, thereafter permitting said material to harden *in*
4 *situ*, the improvements in said method comprising utilizing as the predominant component
5 of said filler, a fused-fibrous material comprised of:
6 from about 1% to about 50% by weight alumina;
7 from about 50% to about 98% by weight silica; and
8 from about 1% to about 5% by weight boron.
- 1 2. The method of claim 1 wherein the filler is comprised of from about 15% to
2 about 30% by weight alumina and from about 65% to about 85% silica.
- 1 3. The method of claim 1 wherein the fused-fibrous material has mean pore diame-
2 ters of greater than 10 microns.
- 1 4. The method of claim 1 wherein the fused-fibrous material is greater than 60% by
2 volume void space.
- 1 5. The method of claim 1 wherein the fused-fibrous material is in particle form.
- 1 6. The method of claim 5 wherein the particles have diameters of about 150
2 microns to about 350 microns.
- 1 7. The method of claim 6 further comprising particles having diameters less than or
2 equal to 5 microns.
- 1 8. A fused-fibrous dental restorative composition comprising:
2 a binder; and
3 a filler, said filler comprised of about 1% to about 50% by weight alumi-
4 na, from about 50% to about 98% percent by weight silica, and boron.
- 1 9. The composition of claim 8 further comprising silicon carbide.

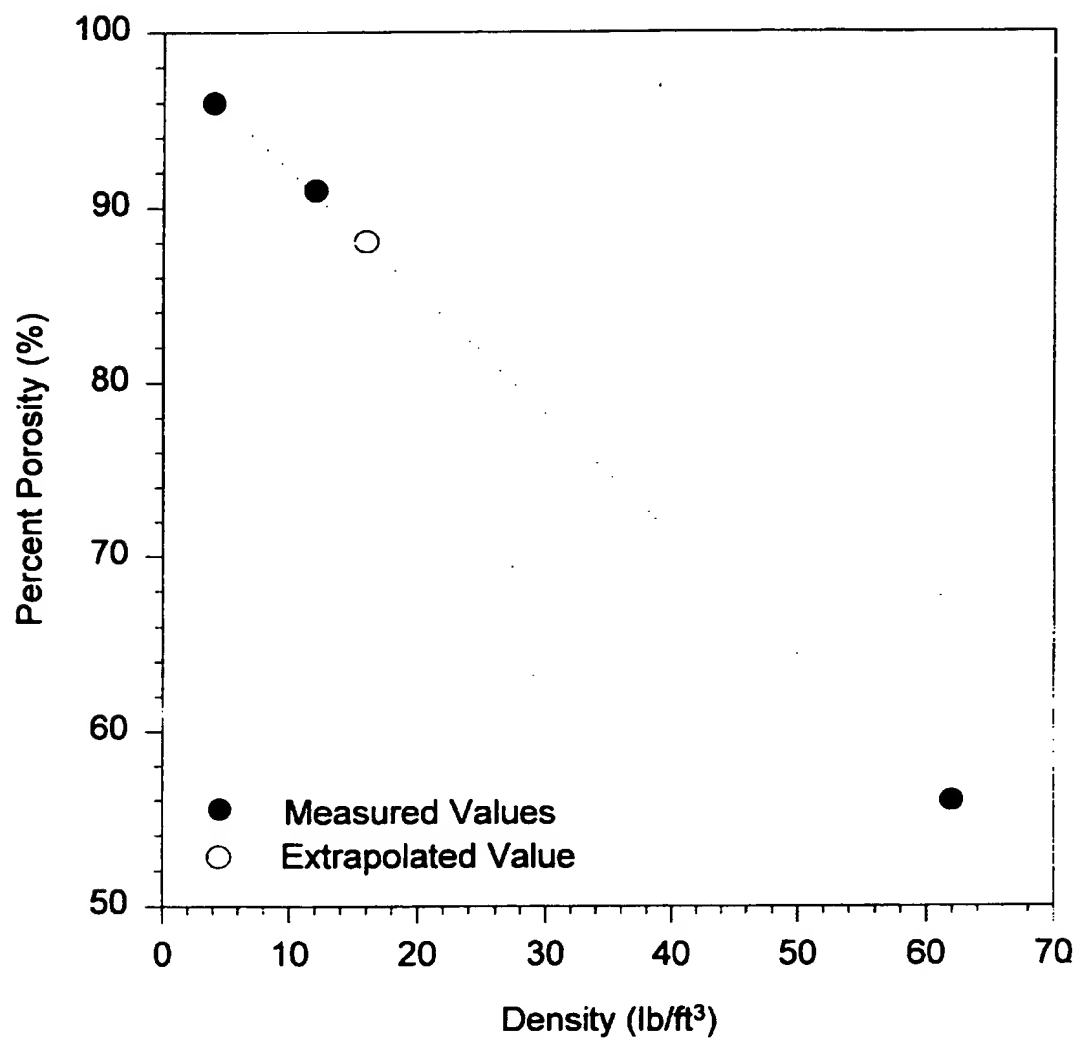
- 1 10. The composition of claim 8 wherein the filler is comprised of about 15% to
2 about 30% by weight alumina and about 65% to about 85% silica.
- 1 11. The composition of claim 8 wherein the filler is comprised of about 21% by
2 weight alumina and about 74% by weight silica.
- 1 12. A method for the filling of a cavity in a tooth comprising:
2 (a) filling said cavity with a settable filling material comprising a binder and
3 filler, wherein said filler comprises at least 60% by volume void space
4 and has mean pore diameters of greater than 10 microns and
5 (b) thereafter allowing said material to harden.
- 1 13. The method of claim 12 wherein the filler is comprised of alumina and silica.
- 1 14. The method of claim 13 wherein the filler further comprises boron.
- 1 15. A fused-fibrous ceramic dental restorative composition manufactured from:
2 alumina fibers, silica fibers and from about 1% to about 5% boron
3 nitride.
- 1 16. The composition of claim 15 further manufactured from silicon carbide.
- 1 17. The composition of claim 15 manufactured from about 2.85% by weight boron
2 nitride.
- 1 18. The composition of claim 15 manufactured from about 21% by weight alumina.
- 1 19. The composition of claim 15 manufactured from about 74% by weight silica.
- 1 20. The composition of claim 15 manufactured from alumina fibers having an average
2 diameter of from about 1 to about 15 microns.

- 1 21. The composition of claim 15 manufactured from silica fibers having an average
2 diameter of from about 1 to about 6 microns.
- 1 22. A fused-fibrous dental restorative composition comprising:
2 alumina; and
3 silica; whereby said composition is greater than about 60%
4 by volume void space and has mean pore diameters greater
5 than about 10 microns.
- 1 23. The composition of claim 22 comprising from about 1% to about 50% by weight
2 alumina.
- 1 24. The composition of claim 22 comprising from about 50% to about 98% by
2 weight silica.
- 1 25. A porous reinforcement for dental restoration materials comprising:
2 a rigid three-dimensional network of fibers fused together at
3 their points of contact wherein said network has mean pore
4 diameters greater than 10 microns.
- 1 26. The reinforcement of claim 25 wherein the network is greater than 60% by
2 volume void space.
- 1 27. The reinforcement of claim 25 wherein the fibers are inorganic.
- 1 28. The reinforcement of claim 25 wherein the fibers are organic.
- 1 29. The reinforcement of claim 25 wherein the fibers are silanated.
- 1 30. A porous reinforcement for dental restoration materials comprising:
2 a rigid three-dimensional network of fibers fused together at
3 their points of contact wherein said network is greater than
4 60% by volume void space.

- 1 31. The reinforcement of claim 30 wherein the fibers are inorganic.
- 1 32. The reinforcement of claim 30 wherein the fibers are organic.
- 1 33. The reinforcement of claim 30 wherein the reinforcement has mean pore diameters
2 greater than about 10 microns.
- 1 34. The reinforcement of claim 30 wherein the fibers are silanated.





PRIMM™ Porosity as a function of Density**Figure 3**

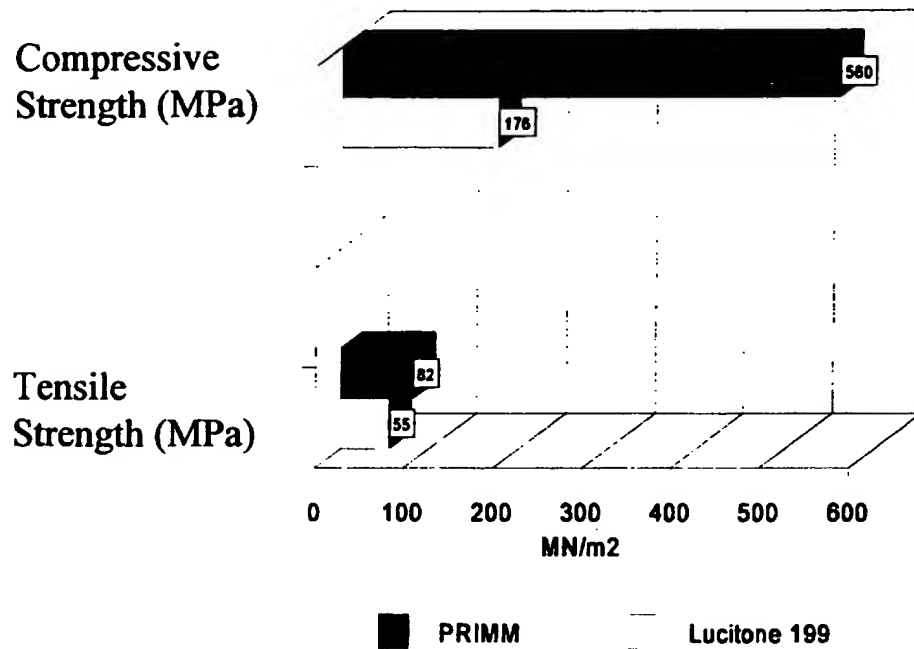


Figure 4

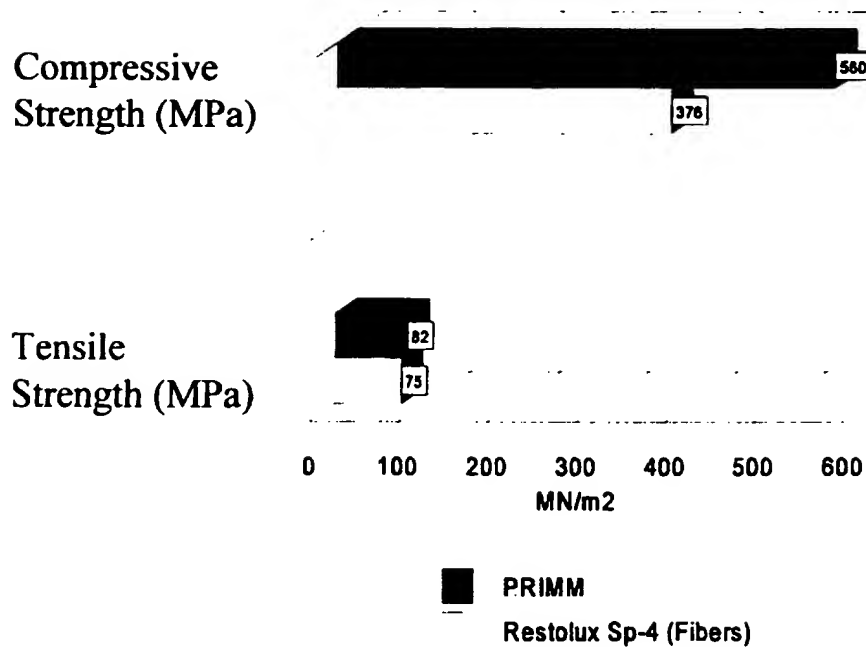


Figure 5

Compressive
Strength (MPa)

Tensile
Strength (MPa)

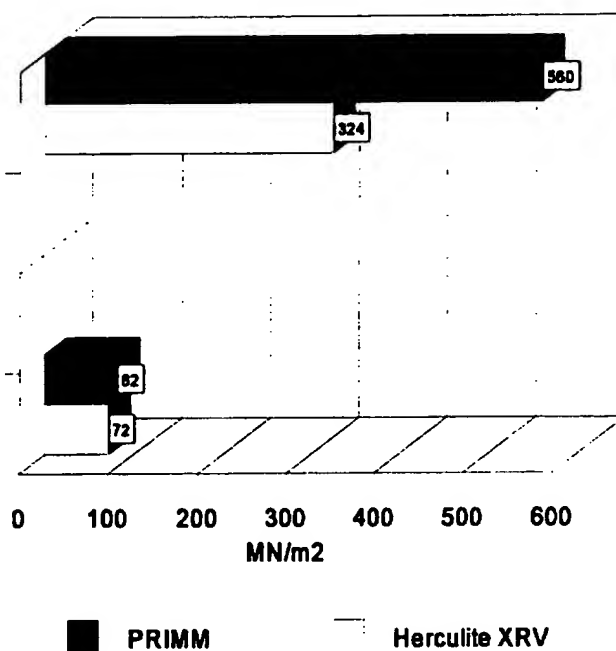


Figure 6

Compressive
Strength (MPa)

Tensile
Strength (MPa)

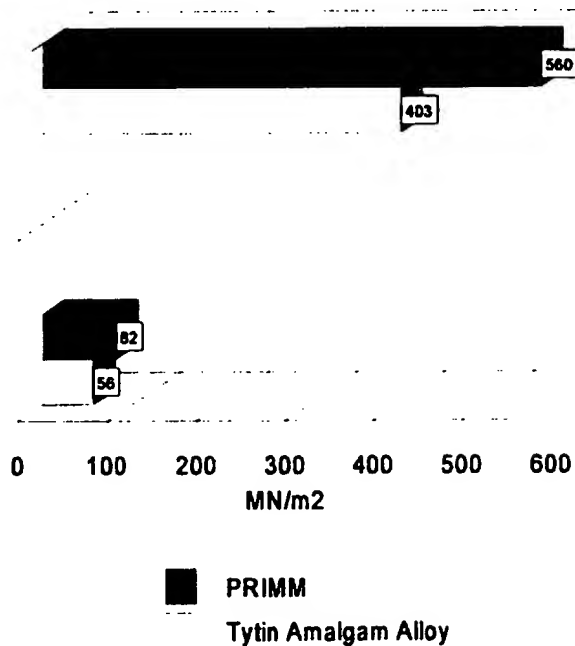


Figure 7

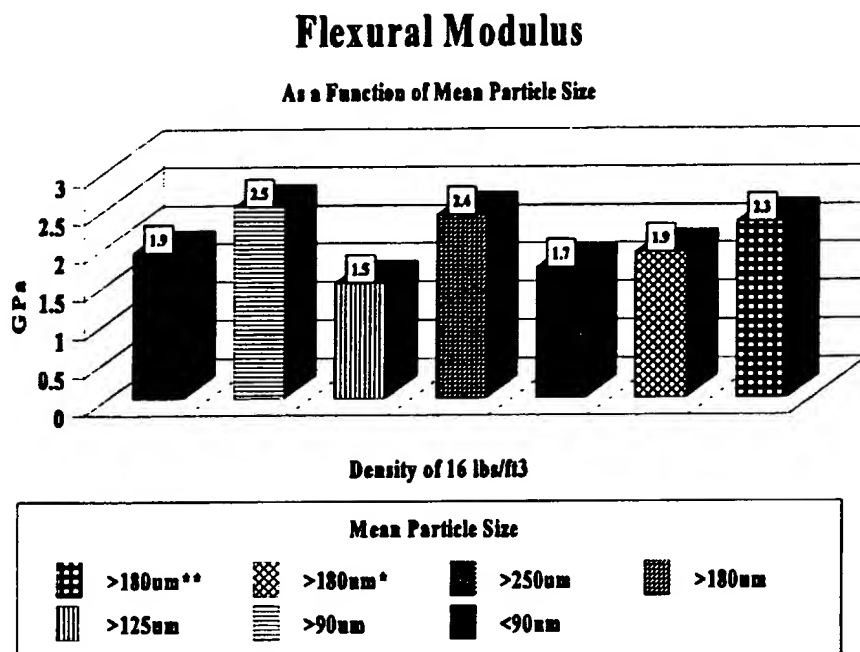


Figure 8

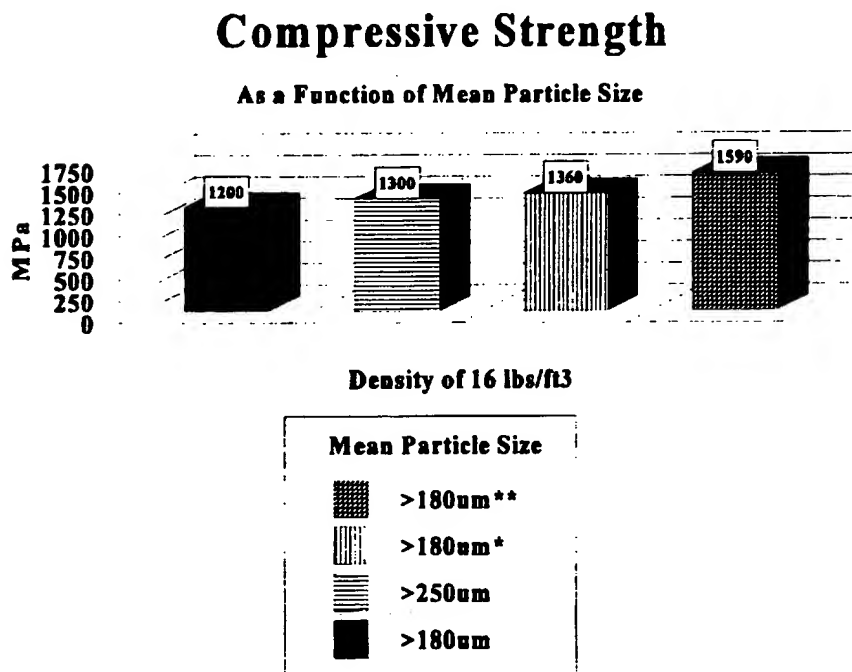


Figure 9

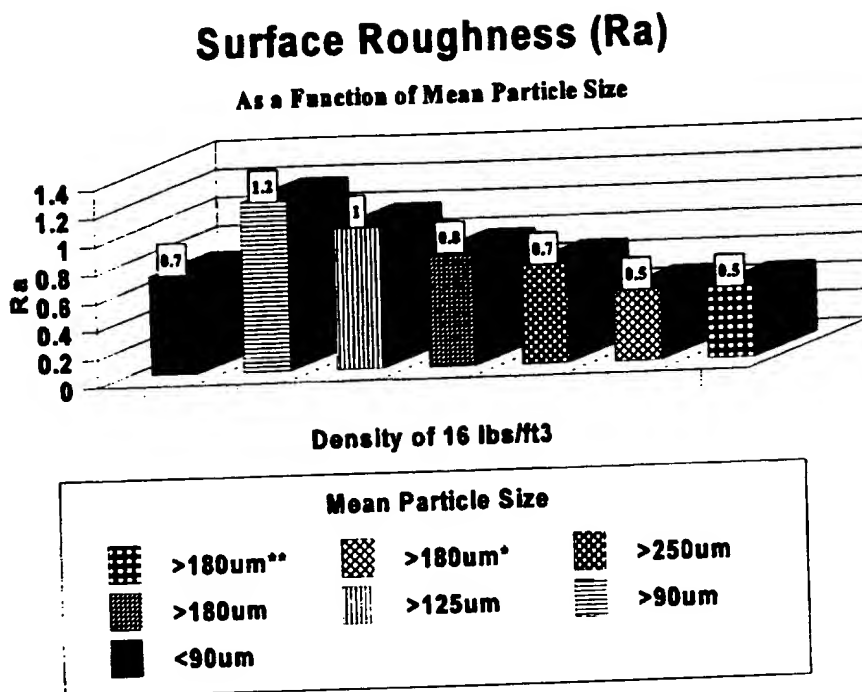


Figure 10

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US96/01800

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : C08K 3/38, 3/08, 3/40; A16F 2/00

US CL : 523/115, 116; 524/404, 441, 492; 433/226, 228.1

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 523/115, 116; 524/404, 441, 492; 433/226, 228.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US, A, 5,273,559 (HAMMAR ET AL) 28 December 1993. See column 6, lines 25-48.	1-34
Y	US, A, 4,650,847 (OMURA ET AL) 17 March 1987. See columns 7-25 and 30-31.	1-34

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	* T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
* A* document defining the general state of the art which is not considered to be part of particular relevance	* X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
* E* earlier document published on or after the international filing date	* Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
* L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	* A* document member of the same patent family
* O* document referring to an oral disclosure, use, exhibition or other means	
* P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

13 APRIL 1996

Date of mailing of the international search report

20 MAY 1996

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